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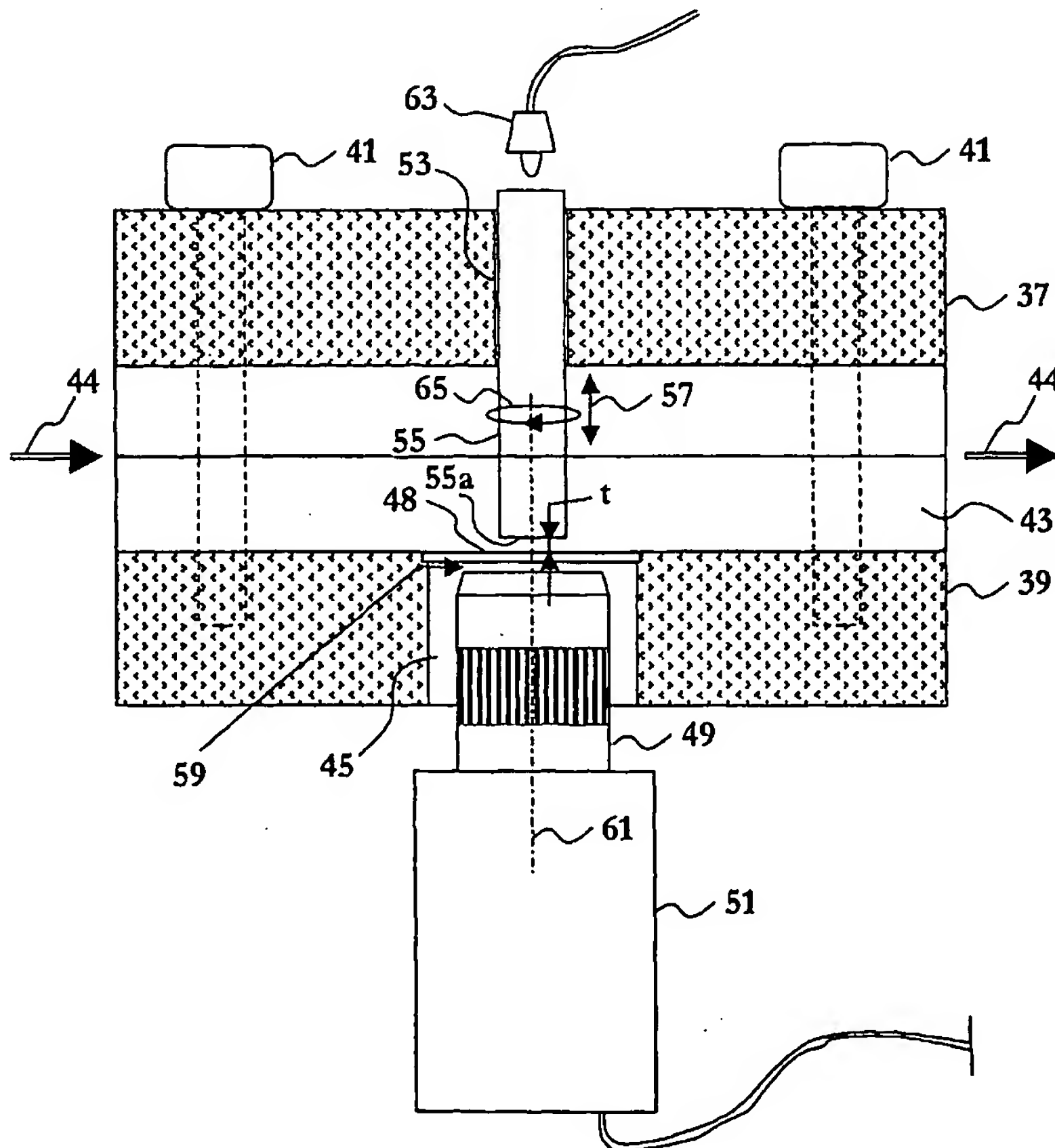
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(54) Title: METHOD AND APPARATUS FOR COUNTING SOMATIC CELLS OR FAT DROPLETS IN MILK



(57) Abstract: A method for counting somatic cells or fat droplets in milk on-line during milking by an automated or semi-automated milking system comprising the steps of: flowing milk as milked by the milking system through a measuring chamber (59); illuminating milk that flows through the measuring chamber; and recording multiple two-dimensional digital images of illuminated milk that flows through the measuring chamber, wherein the images are recorded through a lens system (49) to preferably obtain a spatial resolution better than about 5 microns in the images. Finally, a somatic cell or fat droplet count score of the milk is determined from the images by means of digital image processing, preferably including the use of neural networks.

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## **METHOD AND APPARATUS FOR COUNTING SOMATIC CELLS OR FAT DROPLETS IN MILK**

### **TECHNICAL FIELD OF THE INVENTION**

The present invention relates generally to dairy farming, and  
5 more specifically to methods and apparatuses for counting  
somatic cells or fat droplets in milk.

### **DESCRIPTION OF RELATED ART AND BACKGROUND OF THE INVENTION**

A major cause of loss in dairy farming is an infection, known as  
mastitis, which occurs in an animal's udder. Mastitis is caused  
10 by contagious pathogens invading the udder and producing toxins  
that are harmful to the mammary glands. Generally, mastitis  
starts in one udder quarter.

Somatic cells, predominantly white cells and epithelial cells,  
enter the mammary gland as a result of damage to the alveolar  
15 lining by infection or chemical irritation. The counting of  
somatic cells excreted in the milk has become a widely used  
measure of mammary gland inflammation. The somatic cells can be  
counted by laborious direct microscopic method on stained milk  
smears, or the cell numbers can also be estimated by direct  
20 chemical tests. Other methods measure milk somatic cells  
indirectly or by determining the concentration of various by-  
products of the inflammatory response.

Somatic cell count (SCC), which is the number of white cells per  
milliliter of milk, increases in the bulk tank as mastitis  
25 spreads in the herd. SCC scores are used as an international  
standard in determining milk's quality and price. Most marketing  
organizations and regional authorities regularly measure SCC on  
bulk tank milk and use these scores for penalty deductions

and/or incentive payments. High SCC scores indicate the presence of mastitis in the herd, which is reflected in the average score of the bulk tank. The bulk tank SCC is a good indicator of overall udder health and as good means for evaluating the mastitis control program.

It is also a high correlation between the bulk milk SCC and the average of individual animal counts. It is not uncommon for a few problem animals to be responsible for greater than 50% of the somatic cells in the bulk tank, particularly in small herds. It should be noted that animals with high milk production and intermediate SCC levels can have a significantly higher percentage of SCC contribution to the tank score than some high SCC cows with low production. For high quality milk the SCC should be less than 200,000 cells/ml. Acceptable milk has SCC scores from 200,000 to 500,000 cells/ml. For infected animals, milk SCC scores are between 600,000 and 1.2 million cells/ml.

When an animal in the herd becomes infected with infectious pathogens a rapid drop in milk production will be noted within two to three days. A high level of bacteria in an animal causes an increased level of somatic cells in milk. An increased level of somatic cells in milk results in poorer quality milk products, which are harder to process. The prevention procedures at milking are less efficient especially when the mastitis is in a subclinical phase and there are no visible signs of the disease. Special efforts have to be made at each milking to detect subclinical mastitis in individual animals.

SCC may be measured by CMT (California Mastitis Test) by utilizing the difference in the extent of aggregation reaction depending on the number of somatic cells, when a surfactant is added to the milk. Since a BTB reagent is also included for pH measurement, it is used as an evaluation index for mastitis by

utilizing the fact that increased vascular permeability and accelerated conflict between leukocytes and bacteria during mastitis results in increased salts such as sodium chloride and potassium chloride in the milk, creating a higher alkalinity, and causing a color change from yellow to green and then to blue. The advantages of this measurement are that it can be easily performed by anyone, it can generally distinguish between the presence and absence of mastitis, and it is an extremely low-cost method. The drawbacks of CMT are that diagnosis is difficult until the reaction has occurred, involving the conflict between leukocytes and the bacteria, or after promotion of vascular permeability, and that diagnosis depends on subjective human judgment, so that this method can only serve as an approximate diagnosis method. Diagnosis has been particularly rough in cases where the milk somatic cell count is 300,000/ml or less. The method is thus not suitable to be automated.

Measuring CL (chemiluminescence) activity has also been used for determining the SCC, see e.g. US Pat. No. 6,297,045. A related method is to add to the milk a fluorescent additive, which is absorbed by the cells. By illuminating the milk with light of a particular wavelength the cells will emit a fluorescent light of another characteristic wavelength. By a suitable filter, which filters out light of the characteristic wavelength, the number of cells can be counted.

Such an approach requires that milk samples are taken, that a suitable amount of fluorescent additive has to be added and mixed with the milk, and that particular light sources and filters are used. This is a labor intense and costly procedure. If the method is automated in a milking robot system, particular provisions have to be taken in order to obtain and separate



small amounts of milk, which is representative of the milk from a cow or an udder of a cow.

Mastitis may alternatively be detected by measuring changes in the electrical conductivity of milk as generally, ion  
5 concentration, and thus electrical conductivity, in mastitic milk is higher than in normal milk. Electrical conductivity is generally measured with a DC or AC circuit having a probe positioned in the flow of milk. The most sensitive part of this on-line method is the probe. The probe generally includes two  
10 electrodes to which an AC or DC current is supplied to create an electrical circuit through the milk. The conductivity of the milk is evaluated by measuring the current variations in the circuitry that includes the probe. However, the readings are often inaccurate due to deposits of colloidal materials from the  
15 milk on the electrodes, and also due to polarization. Polarization occurs because some of the ions migrating towards the electrodes are not neutralized and consequently, an offset, or leakage current is generated between the electrodes. The presence of the leakage current results in inaccurate  
20 conductivity readings. Different aspects on milk conductivity measurements have been patented, see e.g. U.S. Pat. Nos. 3,762,371; 5,416,417; 5,302,903; 6,307,362 B1; and 6,378,455 B1.

Conductometry has disadvantages in that it depends on changes occurring by inflammation reaction after the bacteria invade and  
25 conflict with the leukocytes, and therefore it is unsuitable for diagnosis in the initial stages of mastitis, while it has poor reproducibility due to substantial differences in electrolyte components and concentrations in different teats or different cows even with normal milk, such that diagnosis is risky by this  
30 diagnostic method alone.

Another potential problem using milk conductivity measurements to discover mastitis is that the conductivity of the milk is heavily dependent on the milking intervals, see *Influence of different milking intervals on electrical conductivity before alveolar milk ejection in cows*, K. Barth and H. Worstorff, *Milchwissenschaft* 55(7), 2000, p. 363. Thus, the milking intervals have to be taken into consideration if milking times are not as fixed as in conventional milking systems.

#### SUMMARY OF THE INVENTION

10 A general object of the present invention is to provide a method and an apparatus, respectively, for counting somatic cells or fat droplets in milk on-line during milking by an automated milking system, which lack the drawbacks and limitations associated with the prior art described above.

15 A particular object of the invention is to provide such a method and such an apparatus, which are completely automatic and provide a reliable somatic cell or fat droplet count score.

It is a further object of the invention to provide such a method and such an apparatus, which count somatic cells or fat  
20 droplets directly in a milk line of the automated milking system.

It is yet a further object of the invention to provide such a method and such an apparatus, which are capable of providing a separate somatic cell count or fat droplet score for each udder  
25 quarter of a cow.

It is still a further object of the invention to provide such a method and such an apparatus, which are reliable, flexible, of fairly low cost, and relatively easy to implement.

These objects, among others, are according to the present invention attained by methods and apparatuses as specified in the appended patent claims.

Further characteristics of the invention, and advantages thereof, will be evident from the following detailed description of preferred embodiments of the present invention given hereinafter and the accompanying Figs. 1-6, which are given by way of illustration only, and thus are not limitative of the present invention.

#### 10 BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 illustrates schematically, in a perspective view, main components of a milking robot provided with an apparatus for counting somatic cells or fat droplets in milk on-line during milking according to a general embodiment of the present invention.

Figs. 2-3 illustrates schematically, in cross-sectional top and end views, an apparatus for counting somatic cells or fat droplets in milk on-line according to a particular embodiment of the present invention.

20 Figs. 4-6 show three examples of two-dimensional digital images as recorded by the apparatus of Figs. 2-3 during counting of somatic cells or fat droplets.

#### DETAILED DESCRIPTION OF EMBODIMENTS

Fig. 1 illustrates some of the main components of a milking robot. The milking robot comprises four teat cups 11, of which only one is illustrated for sake of simplicity. Each teat cup 11 is connected to a respective milk tube 13, which in turn is connected to an end unit 15 via a respective valve or regulator



17, a respective milk conduit 18, a respective flow meter 19, and a common milk meter 21. The end unit is connected to a vacuum source (not illustrated) via a milk/air separator and a vacuum supply conduit 23.

5 During milking of the teats of a milking animal, the teat cups are attached to the teats of a cow, typically by a robot arm (not illustrated), and vacuum is supplied to the end unit 15 via the vacuum supply conduit 23 to draw milk from the teats of the cow, through the milk lines 13 and into the end unit 15. The  
10 valves or regulators 17 may be used to control the individual vacuum levels in the teat cups 11. The milk from each udder quarter of the cow is measured individually by the flow meters 19, wherafter the weight of the milk from the cow is measured by the common milk meter 21. Finally, the milk is collected in the  
15 end unit 15 and the air is sucked out through the conduit 23.

Further, the milking robot comprises a pump and regulator system 27 for pumping the milk to e.g. a larger milk storage tank (not illustrated) via one 29 of a plurality of milk output lines 29, 31 connected to the end unit 15. Another milk output line 31 may  
20 be used for discarding milk from the milking of a cow, for pumping the milk to another tank (not illustrated), or for pumping the milk to a feed device for feeding calves.

The milking robot is advantageously connected to a computer-based processing and control device 35, which is responsible for  
25 processing and controlling of the milking robot, and comprises typically a microcomputer, suitable software, and a database including information of each of the cows milked by the milking robot, such as e.g. when the respective cow was milked last time, when she was fed last time, her milk production, her  
30 health, etc.

For the purpose of identifying cows, which have an increased SCC scores, e.g. in order to treat or monitor these cows, or in order to direct the milk from them to not mix it with milk from healthy cows or cows having low SCC scores, the present invention presents an improved technique for counting somatic cells in milk on-line during milking.

An inventive apparatus for counting somatic cells or fat droplets in milk on-line during milking, schematically shown by reference numeral 33 in Fig. 1, comprises generally a flat or shallow measuring chamber, a light source, a two-dimensional camera system including a lens system, preferably a microscope, and a digital image processing system. A particular embodiment of the apparatus is illustrated in detail in Figs. 2-3, which embodiment will be described further below.

The flat measuring chamber is arranged so that at least a portion of the milk drawn from the teats of a cow, through the milk lines 13, and into the end unit 15 is flowed through the measuring chamber. The light source is set to illuminate milk that flows through the flat measuring chamber, and the two-dimensional camera system is adapted to repeatedly record two-dimensional digital images of the illuminated milk that flows through the flat measuring chamber. The camera array and the lens system are adapted so that a rather small image area is recorded, but with high magnification. A spatial resolution better than about 5 microns in the two-dimensional digital images is preferred. Finally, the digital image processing system is adapted to determine, e.g. by use of neural networks, a somatic cell or fat droplet count score from the two-dimensional images.

Preferably, the digital image processing system is implemented in the processing and control device 35.

The flat measuring chamber may be arranged in a separate conduit, provided for leading away a portion of the milk from one or several of the milk conduits 18. Optionally, the milk is brought back to the milk conduit(s) 18 or is brought to the end  
5 unit 15 after having passed the flat measuring chamber. Advantageously, however, the flat measuring chamber is arranged within one of the milk conduits 18.

Such solution is adopted by the particular embodiment of the apparatus as being illustrated in Figs. 2-3. A measuring cell  
10 comprises a top and a bottom cell block 37, 39, which when being attached to each other in a fluid tight manner by means of four bolts 41 or similar form a milk passageway 43 from left to right. The passageway 43 has preferably a circular cross section as illustrated. The measuring cell is mounted in one of  
15 the milk conduits 18 so that milk flows through the passageway 43 as indicated by arrows 44. Alternatively the cell blocks 37, 39 are designed to form a milk passageway of other cross sectional shape, e.g. quadratic or rectangular.

Further, the bottom cell block 39 of the measuring cell is  
20 provided with a substantially vertical through hole 45. The surface of the bottom cell block 39, which together with a corresponding surface of the top cell block 37, form the passageway 43, is shaped to be plane within a major portion of a given area. A light transparent plate 48 fitted within the  
25 flat portion is glued to the bottom cell block 39 in a fluid tight manner. The position of the plane surface portion of the bottom cell block 39 is selected so that the upper surface of the plate 48 is in level with the lowest portion of the surface forming the passageway 43 outside the area. The passageway  
30 surface of the bottom cell block 39 within the given area, but outside the plane surface portion, may be shaped to obtain a

smooth transition to the passageway surface of the bottom cell block 39 outside the given area. By providing smooth surfaces within the measuring cell, pockets where milk may be accumulated are avoided. The size of the hole 45 is selected  
5 such that the front portion of a two-dimensional camera system 51, e.g. CCD-based system, provided with a lens system 49, preferably a microscope, for magnification can be inserted into the hole 45 as illustrated.

The top cell block 37 of the measuring cell is provided with a  
10 substantially vertical through hole 53, preferably smaller than the hole 45, and aligned with the hole 45. A rod 55 is fitted to be inserted to the through hole 53 so that a flat end surface 55a of the rod 55 is located in the passageway 43 opposite to and parallel with the plate 48. The rod 55 is  
15 tightly fitted in the through hole 53 to prevent milk from leaking out through the hole 53, and is movable in a vertical direction as is indicated by arrow 57.

The flat measuring chamber 59 is defined as the space between the plate 48 and the flat end surface 55a of the rod 55. Thus,  
20 the flat measuring chamber 59 is open in directions being parallel with the plate 48 and the surface 55a, and orthogonal to a general direction of the flow of milk as indicated by the arrows 44. During SCC measurements the thickness  $t$  of the measuring chamber 59, i.e. the dimension of the measuring  
25 chamber 59 in a direction parallel with the optical axis 61 of the camera system 51 during measurements, is preferably smaller than about 100 microns, more preferably smaller than about 50 microns, and most preferably smaller than about 10 microns. It is important to obtain a depth of field and focusing of the  
30 camera system 51 so that the images are sharp; and to reduce

the probability of cells "hiding" behind an imaged cell. Such cells will obviously not be counted.

The rod 55 is preferably light transparent to allow for illumination of the milk that flows through the flat measuring chamber 59 by a light source, schematically indicated by 63, through the rod 55. It shall, however be appreciated by the man skilled in the art that other illumination techniques may be used including i.a. mirror and beamsplitter arrangements. Milk in the flat measuring chamber 59 may be illuminated from above as illustrated or from below, i.e. from the camera system 51 side. In the latter instance the end surface 55a of the rod 55 may be light reflecting.

In general, light as transmitted through milk in the measuring chamber is recorded by the camera system. Alternatively or additionally, light as reflected by milk in the measuring chamber is recorded. Further, the orientation of the measuring chamber 59 and the camera system 51 may differ from what is illustrated in Figs. 2-3.

The milk is sucked through the lines 18 intermittently and is mixed with air. Thus, it is particularly advantageous to have the measuring chamber 59 arranged at the very bottom of the passageway 43 as it is most probable that milk will pass through there due to gravity. In order to assure that milk is not clogged in the measuring chamber 59, the rod may be rotated around the axis 61 continuously during measurements as being indicated by arrow 65. The rod may be moved vertically and be rotated automatically by means of a motor (not illustrated) connected to the processing and control device 35.

The camera system 51 is preferably provided with a microscope or tele/macro photo lens system 49 to record strongly magnified



two-dimensional images. Preferably, the camera system 51 provides for a spatial resolution in the two-dimensional digital images better than 2 microns, more preferably better than about 1 micron, and most preferably better than about 0.5 5 microns. As a result thereof very small areas are recorded and probably a very large number of images have to be recorded in order to provide accurate and precise SCC scores.

Three different two-dimensional images as recorded by a SCC measuring apparatus according to the principles of the present 10 invention, but set up in a laboratory environment, are illustrated in Figs. 4-6.

In the first image (Fig. 4) only fat droplets are visible, whereas in the second and third images (Figs. 5-6) several somatic cells are identified among a large number of fat 15 droplets (the somatic cells are indicated by the arrows). As can be seen in Figs. 5-6 the somatic cells look quite different than the fat droplets and these differences are used by the digital image processing system to distinguish the different particles in the images. Generally, the digital image 20 processing employed includes the analysis of number, size, shape, structure, morphological structure, density and/or composition of particles found in each image as revealed by the reflection and/or transmission properties of the particles as found in the images recorded. Preferably, the image processing 25 system uses neural networks.

Using a 600x400 pixel CCD-camera provided with a microscope to record images covering an area of  $0.3 \times 0.2 \text{ mm}^2$  the spatial resolution in the images is estimated to be about 0.5 microns. Using a measuring chamber with a thickness of about 0.1 mm each 30 sample volume imaged amounts to  $0.6 \times 10^{-6} \text{ ml}$ . Thus given a SCC score of 1 million cells/ml, which may be a typical score for



an infected cow an average of only 0.6 cells/image will be found in each image. By recording a large number, e.g. thousands, of images, and by means of digital image processing of these images a somatic cell count score can be determined.

- 5 The somatic cells are in some instances, e.g. when the milk is mastitic, predominantly white cells, and thus the somatic cell count score may be a count score of white cells. In other instances, e.g. for healthy animals having naturally high SCC scores, the number of epithelial cells are higher. In still  
10 other instances, e.g. in case of serious disease or injury, the number of red blood cells may be estimated.

From number and size of fat droplets in the images a content of fat may also be estimated using digital imaging processing.

- While the particular embodiment of the SCC measuring apparatus  
15 has been described as being mounted in one of the milk conduits 18, and thus measures SCC in a single udder quarter, it may be connected downstream of the point where milk from the udder quarters are mixed. For instance in a milking machine where the teat cups are connected to a single milk line via a claw  
20 (upstream of the end unit), the SCC measuring apparatus may be located in this single milk line. However, since mastitis often starts in one or maybe two udder quarters, this is not the most preferred solution as the detection sensitivity for mastitis is reduced when milk from infected udder quarters are mixed with  
25 milk from healthy udder quarters before the SCC measurement takes place.

- The most flexible solution is to have a measuring cell mounted in each one of the milk lines 18 in the robot of Fig. 1, and then to provide one light source and one camera system for each  
30 measuring cell, or to provide a single light source or camera

system which is alternately used for SCC measurement of milk that is flowed through the various measuring cells. Then, the SCC scores for the different udder quarters may be compared to obtain a very sensitive detection of mastitis or increased SCC  
5 scores in milk from individual udder quarters.

It shall further be appreciated, that by implementing the above-identified flexible solution, a milking robot with four end units — one for each udder quarter, milk could be transported and taken care of on an udder quarter individual  
10 basis, e.g. milk from udder quarters having low SCC score is collected in one tank and milk from udder quarters having high SCC score is collected in an other tank.

It shall still further be appreciated by the person skilled in the art that the present invention may be implemented in  
15 virtually any kind of automated or semi-automated milking system.

## CLAIMS

1. A method for counting cells or fat droplets in milk on-line during milking of a milking animal, characterized by the steps of:

- 5 - flowing at least a portion of the milk as obtained during said milking of said milking animal through a measuring chamber (59);
- illuminating milk that flows through said measuring chamber;
- repeatedly recording two-dimensional digital images of  
10 illuminated milk that flows through said measuring chamber, said two-dimensional digital images being recorded through a lens system (49), preferably a microscope; and
- determining a somatic cell or fat droplet count score from said two-dimensional images by means of digital image  
15 processing.

2. The method of claim 1 wherein said at least portion of the milk flowed through said measuring chamber (59) is free from toxic additives.

3. The method of claim 1 wherein said at least portion of the  
20 milk flowed through said measuring chamber (59) is pure natural milk, optionally mixed with air, but free from any chemical additives.

4. The method of any of claims 1-3 wherein said repeatedly recordings of two-dimensional digital images are performed to  
25 obtain a spatial resolution better than about 5 microns, preferably better than about 2 microns, more preferably better

than about 1 micron, and most preferably better than about 0.5 microns, in said two-dimensional digital images.

5. The method of any of claims 1-4 wherein said measuring chamber has a dimension (t) smaller than about 100 microns, preferably smaller than about 50 microns, and more preferably smaller than about 10 microns, in a direction parallel with the optical axis (61) of said lens system during said repeated recordings.

6. The method of any of claims 1-5 wherein said digital image processing includes the analysis of number, shape, size, structure, density and/or composition of particles found in each image as revealed by the reflection and/or transmission properties of the particles recorded spatially resolved by said camera system.

7. The method of any of claims 1-6 wherein said digital image processing includes the use of neural networks.

8. The method of any of claims 1-7 wherein said at least portion of said milk, which is flowed through said measuring chamber, is lead away from a milk line (13) of a milking machine used to collect the milk as obtained during said milking of said milking animal.

9. The method of claim 8 wherein said at least portion of said milk, which is lead away from said milk line, is brought back to said milk line or brought to a milk collecting container, after having been flowed through said measuring chamber.

10. The method of any of claims 1-7 wherein said at least portion of said milk is flowed through said measuring chamber (59) within a milk line (13) of a milking machine used to

collect the milk as obtained during said milking of said milking animal.

11. The method of any of claims 1-10 wherein said milking of said milking animal is performed by an automated or semi-automated milking system, which comprises a plurality of teat cups (11), each of which being connected to a respective milk line (13), which milk lines in turn are connected to a container (15) via a claw and a single milk line, wherein, during milking of the teats of said milking animal, said plurality of teat cups are attached to the teats of the milking animal and vacuum (23) is supplied to said container to draw milk through said milk lines, said claw, said single milk line and into said container.

12. The method of any of claims 1-10 wherein said milking of said milking animal is performed by an automated or semi-automated milking system, which comprises a plurality of teat cups (11), each of which being connected to a respective milk line (13), which milk lines in turn are connected to a container (15) wherein, during milking of the teats of said milking animal, said plurality of teat cups are attached to the teats of the milking animal and vacuum (23) is supplied to said container to draw milk through said milk lines and into said container, wherein said milk is drawn in separate milk lines (13) all the way to said container.

13. The method of any of claims 1-12 wherein said somatic cell or fat droplet count score is a count score of white cells.

14. The method of any of claim 11 or 12 wherein said container is provided with a plurality of milk output lines (29, 31); and said milk drawn through the milk lines and into said container

is output through one of said plurality of milk output lines depending on said somatic cell or fat droplet count score.

15. The method of any of claims 1-14 wherein a content of fat is estimated from said two-dimensional images by means of said  
5 digital imaging processing.

16. The method of claim 15 wherein said content of fat is estimated from number and size of fat droplets in said two-dimensional images.

17. The method of claim 12 wherein

- 10 - a measuring chamber (59) is provided in each milk line;
- at least a portion of the milk drawn through the respective milk lines is passed through the respective measuring chambers;
- milk that flows through the respective measuring chambers is illuminated;
- 15 - two-dimensional digital images of illuminated milk that flows through the respective measuring chambers is repeatedly recorded, where said two-dimensional digital images are recorded through a lens system to obtain a spatial resolution better than about 5 microns in said two-dimensional digital  
20 images; and
- somatic cell or fat droplet count scores for milk drawn through the respective milk lines are determined from said two-dimensional images by means of digital image processing.

18. An apparatus for counting somatic cells or fat droplets in  
25 milk on-line during milking of a milking animal,  
characterized in:



- a measuring chamber (59), through which the milk as obtained during said milking of said milking animal is flowed;
- a light source system (63) for illuminating milk that flows through said measuring chamber;
- 5 - a two-dimensional camera system (51) including a lens system (49), preferably a microscope, for repeatedly recording two-dimensional digital images of illuminated milk that flows through said measuring chamber, where said two-dimensional digital images are recorded through said lens system; and
- 10 - a digital image processing system (35) for determining a somatic cell or fat droplet count score from said two-dimensional images.

19. The apparatus of claim 18 wherein said at least portion of the milk flowed through said measuring chamber (59) is free  
15 from toxic additives.

20. The apparatus of claim 18 wherein said at least portion of the milk flowed through said measuring chamber (59) is pure milk natural milk, optionally mixed with air, but free from any chemical additives.

20 21. The apparatus of any of claims 18-20 wherein said two-dimensional camera system provides for a spatial resolution in said two-dimensional digital images better than about 5 microns, preferably better than about 2 microns, more preferably better than about 1 micron, and most preferably  
25 better than about 0.5 microns.

22. The apparatus of any of claims 18-21 wherein said measuring chamber has a dimension (t) smaller than about 100 microns, preferably smaller than about 50 microns, and more preferably

smaller than about 10 microns, in a direction parallel with the optical axis (61) of said lens system during said repeated recordings.

23. The apparatus of any of claims 18-22 wherein said digital  
5 image processing system is adapted to analyze number, shape, size, structure, density and/or composition of particles found in each image as revealed by reflection and/or transmission properties of the particles as recorded by said camera system.

24. The apparatus of any of claims 18-23 wherein said digital  
10 image processing system is adapted to use neural networks in determining said somatic cell or fat droplet count score from said two-dimensional images.

25. The apparatus of any of claims 18-24 wherein

- said milking of said milking animal is performed by an  
15 automated or semi-automated milking system, which comprises a plurality of teat cups (11), each of which being connected to a respective milk line (13), which milk lines in turn are connected to a container (15), wherein, during milking of the teats of said milking animal, said plurality of teat cups are  
20 attached to the teats of the milking animal and vacuum (23) is supplied to said container to draw milk through said milk lines and into said container; and

- said measuring chamber (59), through which said at least  
portion of said milk is flowed, is arranged within one of said  
25 milk lines (13).

26. The apparatus of claim 25 wherein said measuring chamber is defined by a light transparent plate (48) mounted in a wall of said one of said milk lines, through which said two-dimensional camera system is adapted to record said two-dimensional images;

and an oppositely located substantially flat and parallel surface (55a).

27. The apparatus of claim 26 wherein said measuring chamber is open in directions being parallel with said light transparent plate and said substantially flat surface, and orthogonal to a  
5 general direction of the flow (44) of said at least portion of said milk.

28. The apparatus of claim 26 or 27 wherein said substantially flat surface is rotatable (65) around an axis being orthogonal  
10 to said light transparent plate and said substantially flat surface.

29. The apparatus of any of claims 26-28 wherein said substantially flat surface is an end surface of a rod (55).

30. The apparatus of claim 29 wherein said rod is light  
15 transparent to allow for illumination through said rod of said milk that flows through said measuring chamber.

31. The apparatus of any of claims 25-30 wherein said container is provided with a plurality of milk output lines (29, 31); and said apparatus further comprises a pump and regulator system  
20 (27) connected to said digital image processing system (35) for pumping said milk drawn through the milk lines and into said container out through one of said plurality of milk output lines depending on said somatic cell or fat droplet count score.

25 32. The apparatus of any of claims 25-30 wherein

- each of said milk lines is provided with a measuring chamber, through which a portion of the milk drawn through the respective milk line is passed;

- said light source system is adapted to illuminate milk that flows through each of said measuring chambers;

- said two-dimensional camera system is adapted to repeatedly record two-dimensional digital images of illuminated milk that  
5 flows through each of said measuring chambers; and

- said digital image processing system is adapted to determine a somatic cell or fat droplet count score for milk drawn through each of said milk lines from said two-dimensional images.

10 33. The apparatus of any of claims 18-24 wherein

- said milking of said milking animal is performed by an automated or semi-automated milking system, which comprises a plurality of teat cups (11), each of which being connected to a respective milk line (13), wherein, during milking of the teats  
15 of said milking animal, said plurality of teat cups are attached to the teats of the milking animal and vacuum (23) is supplied to said teat cups through said milk lines to draw milk through said milk lines;

- each of said milk lines is provided with a measuring chamber,  
20 through which a portion of the milk drawn through the respective milk line is passed;

- said light source system is adapted to illuminate milk that flows through each of said measuring chambers;

- said two-dimensional camera system is adapted to repeatedly  
25 record two-dimensional digital images of illuminated milk that flows through each of said measuring chambers;

- said digital image processing system is adapted to determine a somatic cell or fat droplet count score for milk drawn

through each of said milk lines from said two-dimensional images; and

- a directing means connected to said digital image processing system for directing milk drawn through each of the respective milk lines into a selected one of a plurality of containers depending on the respective somatic cell or fat droplet count score.

34. A milking robot comprising the plurality of teat cups (11), the plurality of milk lines (13), the container (15), and the apparatus for counting somatic cells or fat droplets of any of claims 25-33.

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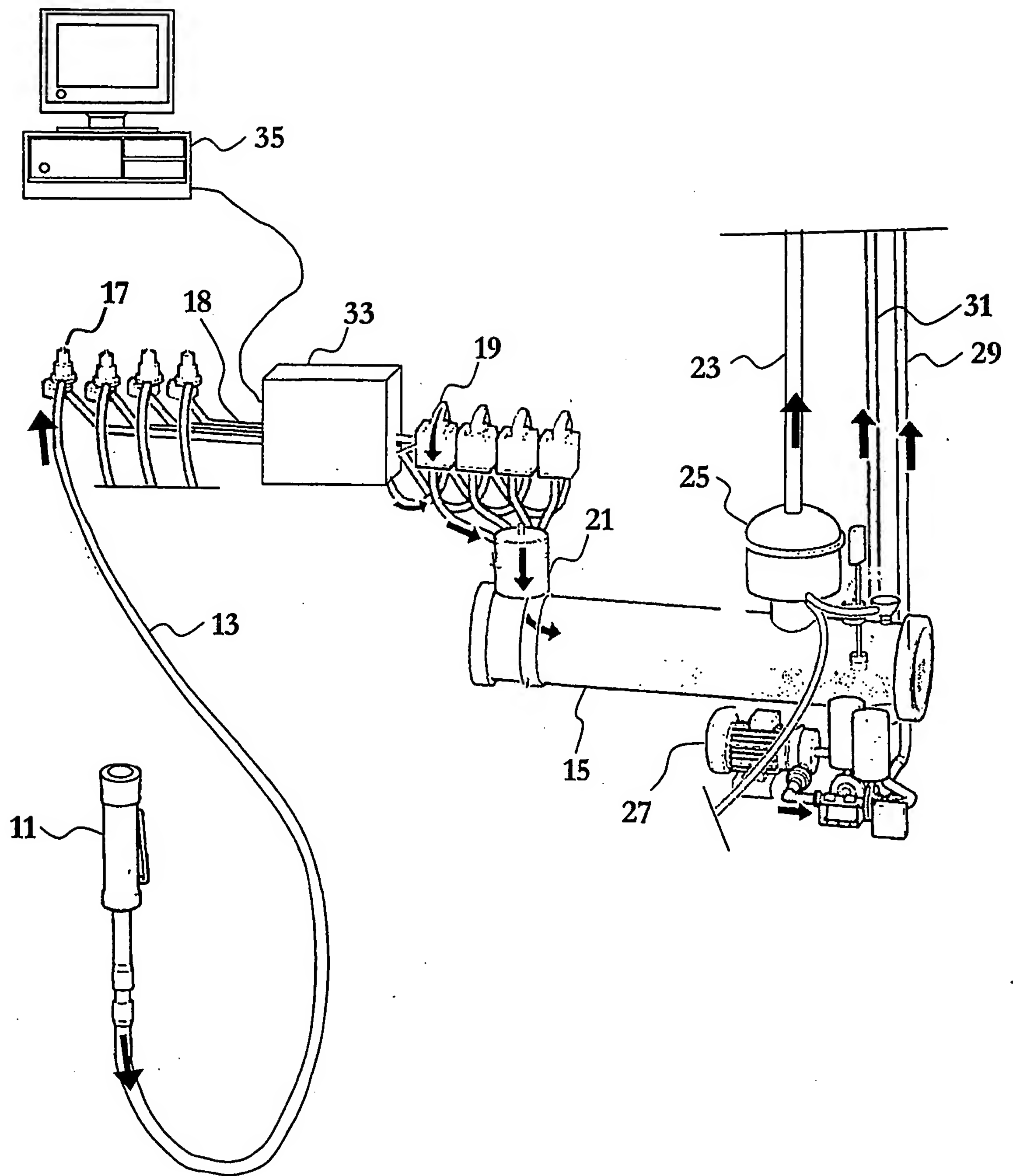


Fig. 1



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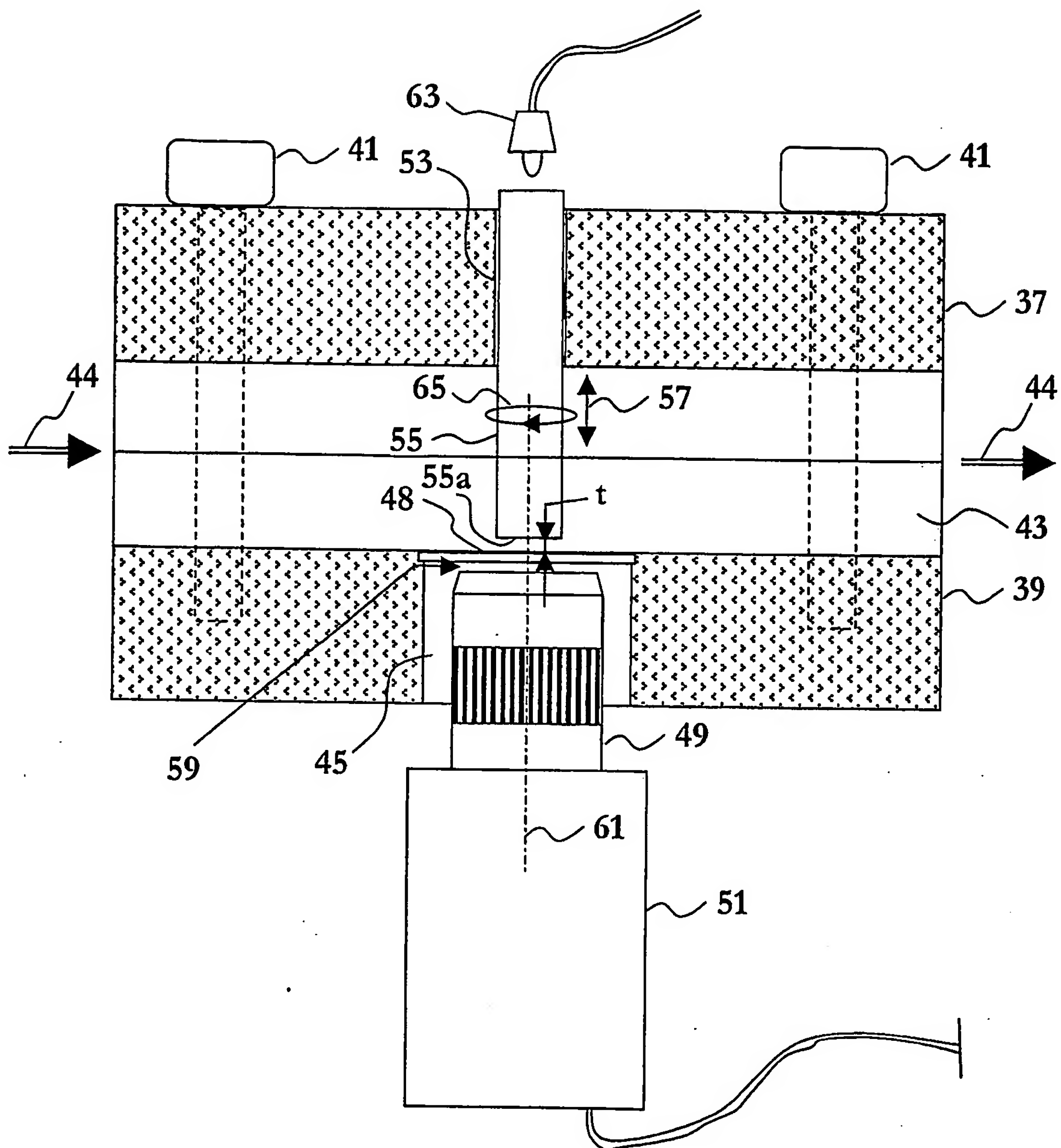


Fig. 2

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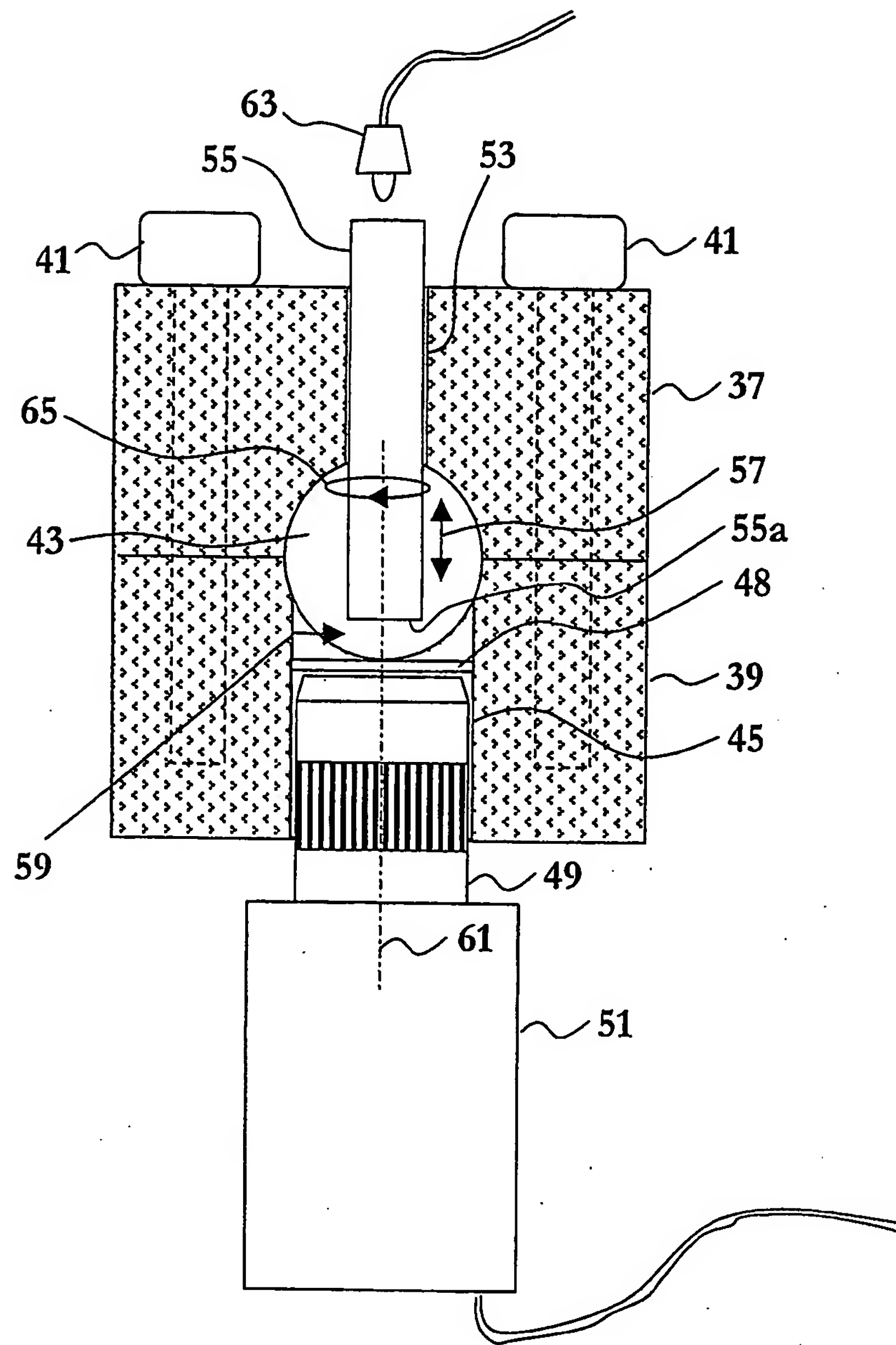


Fig. 3

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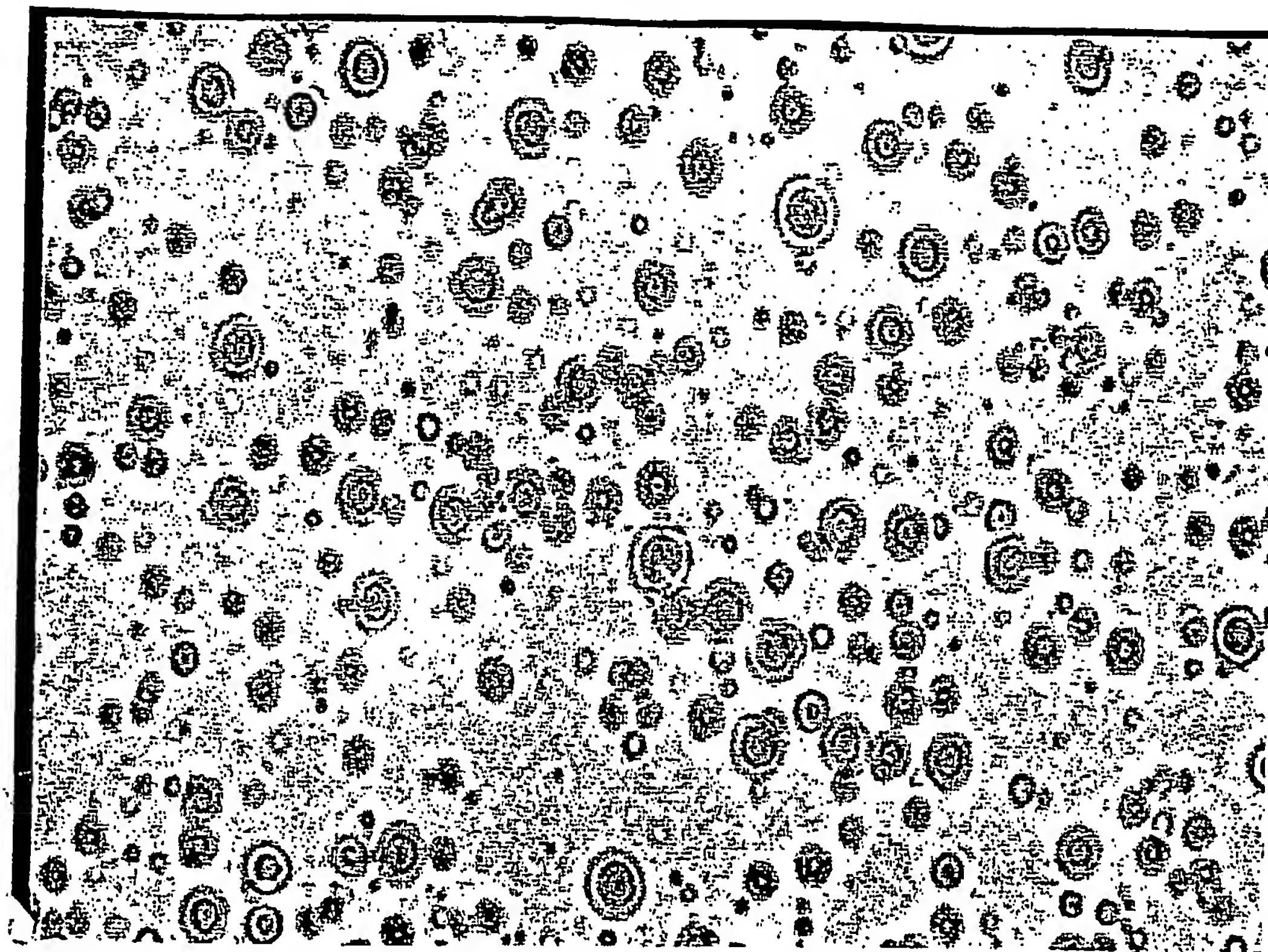


Fig. 4

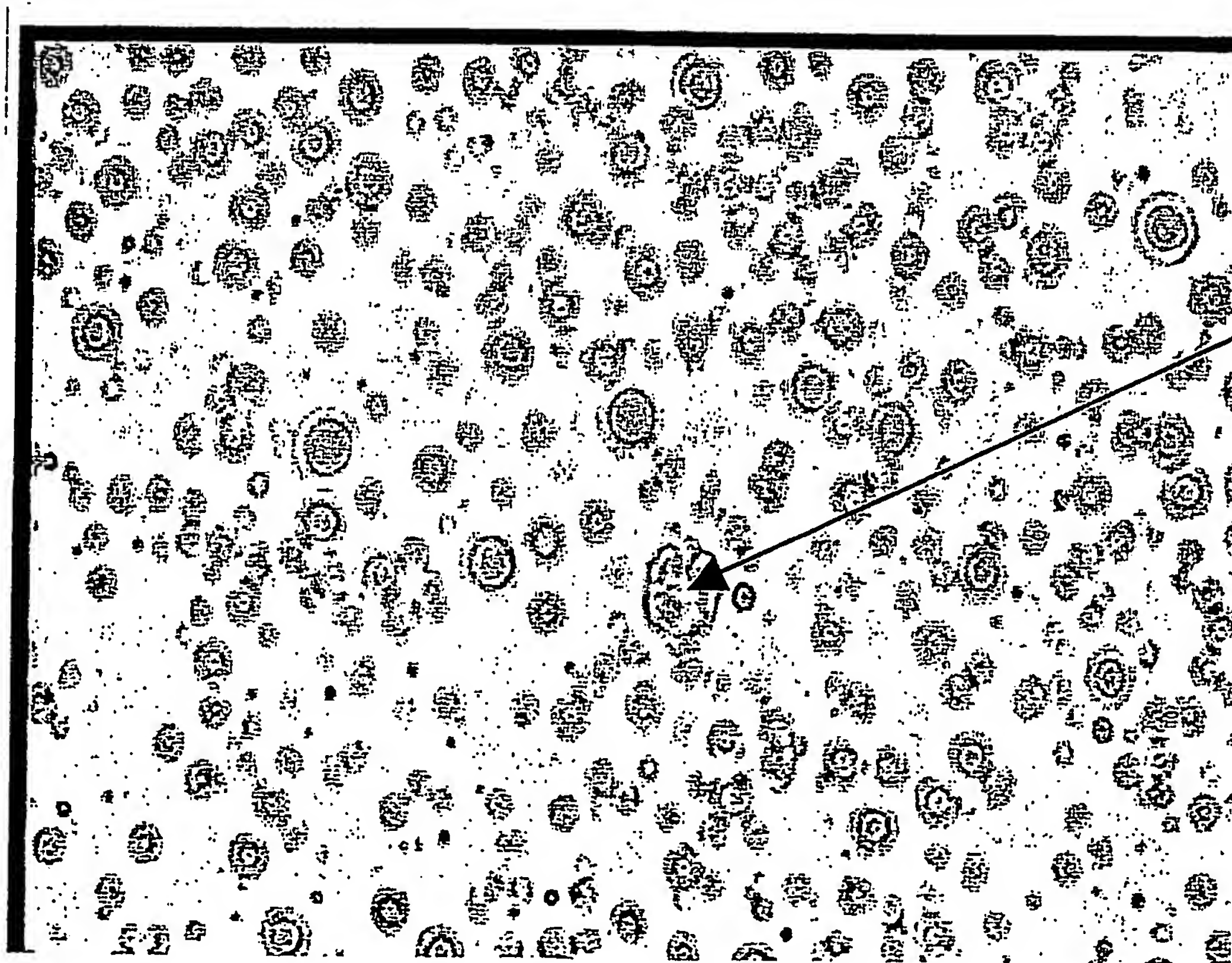


Fig. 5



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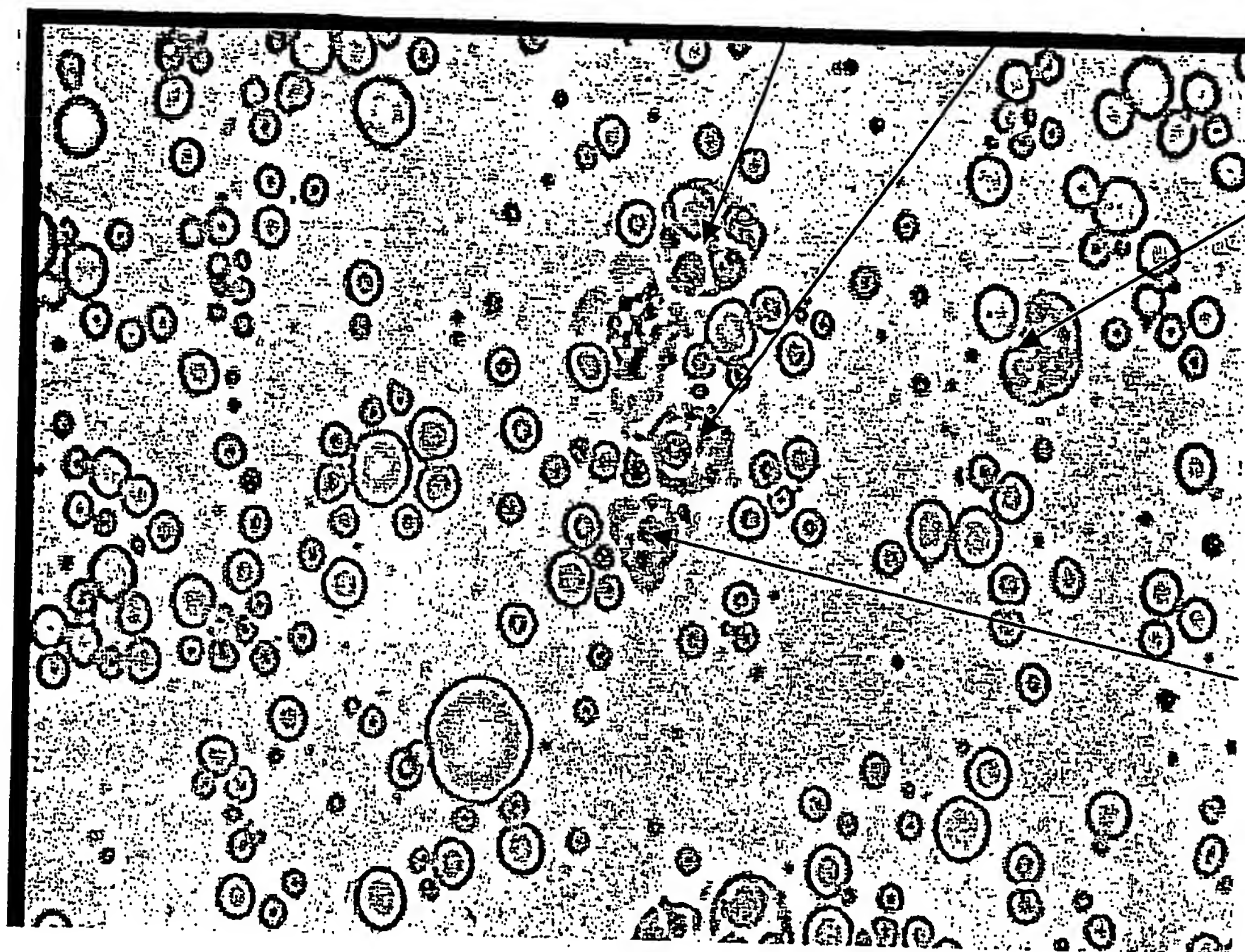


Fig. 6

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 2004/000204

## A. CLASSIFICATION OF SUBJECT MATTER

IPC7: A01J 5/013, G01N 21/01

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: A01J, G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-INTERNAL, WPI DATA, PAJ

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 0027183 A1 (CHEMOMETEC A/S), 18 May 2000 (18.05.2000), page 11, line 13; page 13, line 16 - line 20; page 14, line 9 - line 32, page 15, line 6-line 16 --	1-3,6,11-14, 17-20,23, 31-34
A	EP 1180675 A2 (CHEMOMETEC A/S), 20 February 2002 (20.02.2002), page 2, line 52 - line 54; page 3, line 52 - line 58; page 5, line 18 - line 19 --	1-34
A	EP 1000535 A1 (MAASLAND N V), 17 May 2000 (17.05.2000), column 6, figure 1, abstract --	1-34

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

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"A" document defining the general state of the art which is not considered to be of particular relevance

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"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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"&amp;" document member of the same patent family

Date of the actual completion of the international search

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Date of mailing of the international search report

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## INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 2004/000204

## C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5594544 A (HORIUCHI ET AL), 14 January 1997 (14.01.1997), column 2, line 17 - line 27, figure 1  --	1,4-7,15,17, 18,21-26
A	US 6297505 B1 (FRANDSEN ET AL), 2 October 2001 (02.10.2001), figures 2,4,5, abstract  --	3,9,10,22
A	US 6104483 A (SEBOK ET AL), 15 August 2000 (15.08.2000), column 3, line 35 - line 37, figure 3, abstract  -- -----	1,27,28



## INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 2004/000204

WO	0027183	A1	18/05/2000	AT	228294	T	15/12/2002
				AU	1032100	A	29/05/2000
				CA	2349549	A	18/05/2000
				DE	69904228	D,T	24/12/2003
				DK	1126757	T	30/12/2002
				EP	1126757	A,B	29/08/2001
				SE	1126757	T3	
				JP	2002529057	T	10/09/2002
-----							
EP	1180675	A2	20/02/2002	AT	236387	T	15/04/2003
				AU	737298	B	16/08/2001
				AU	739824	B	18/10/2001
				AU	7205798	A	27/11/1998
				AU	7372298	A	27/11/1998
				CA	2288801	A	12/11/1998
				CA	2288996	A	12/11/1998
				DE	69812928	D,T	04/03/2004
				DK	980516	T	22/04/2003
				EP	0980516	A,B	23/02/2000
				EP	0983378	A	08/03/2000
				EP	1180676	A	20/02/2002
				EP	1180677	A	20/02/2002
				IL	132688	D	00/00/0000
				JP	2001526780	T	18/12/2001
				JP	2002503097	T	29/01/2002
				NZ	500686	A	25/05/2001
				NZ	500687	A	25/05/2001
				US	6710879	B	23/03/2004
				WO	9850577	A	12/11/1998
				WO	9850777	A	12/11/1998
-----							
EP	1000535	A1	17/05/2000	SE	1000535	T3	
				AT	243926	T	15/07/2003
				AU	765969	B	09/10/2003
				AU	5835199	A	18/05/2000
				CA	2288452	A	12/05/2000
				DE	69909215	D	00/00/0000
				DK	1000535	T	27/10/2003
				JP	2000146832	A	26/05/2000
				NL	1010540	C	00/00/0000
				NZ	500916	A	30/11/2001
				US	6197538	B	06/03/2001
-----							
US	5594544	A	14/01/1997	DE	4437758	A	27/04/1995
				JP	7120375	A	12/05/1995
-----							

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 2004/000204

US	6297505	B1	02/10/2001	AT	223041	T	15/09/2002
				AU	731963	B	05/04/2001
				AU	4862897	A	29/05/1998
				CA	2270222	A	14/05/1998
				DE	69715024	D,T	02/01/2003
				DK	122196	A	02/05/1998
				DK	173073	B	20/12/1999
				EP	0935749	A,B	18/08/1999
				JP	2001509254	T	10/07/2001
				KR	2000052993	A	25/08/2000
				NZ	335597	A	28/02/2000
				PL	186428	B	30/01/2004
				PL	333234	A	22/11/1999
				WO	9820338	A	14/05/1998
				AU	737907	B	06/09/2001
				AU	9737598	A	24/05/1999
				CA	2307774	A	14/05/1999
				EP	1031022	A	30/08/2000
				NZ	504777	A	20/12/2002
				US	6573988	B	03/06/2003
				WO	9923473	A	14/05/1999

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US	6104483	A	15/08/2000	NONE
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